

RESEARCH ARTICLE

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Expression of *Shigella flexneri* *gluQ-rs* gene is linked to *dksA* and controlled by a transcriptional terminator

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Abstract

Background: Glutamyl queuosine-tRNA^{Asp} synthetase (GluQ-RS) is a paralog of the catalytic domain of glutamyl-tRNA synthetase and catalyzes the formation of glutamyl-queuosine on the wobble position of tRNA^{Asp}. Here we analyze the transcription of its gene in *Shigella flexneri*, where it is found downstream of *dksA*, which encodes a transcriptional regulator involved in stress responses.

Results: The genomic organization, *dksA-gluQ-rs*, is conserved in more than 40 bacterial species. RT-PCR assays show co-transcription of both genes without a significant change in transcript levels during growth of *S. flexneri*. However, mRNA levels of the intergenic region changed during growth, increasing at stationary phase, indicating an additional level of control over the expression of *gluQ-rs* gene. Transcriptional fusions with *lacZ* as a reporter gene only produced β-galactosidase activity when the constructs included the *dksA* promoter, indicating that *gluQ-rs* do not have a separate promoter. Using bioinformatics, we identified a putative transcriptional terminator between *dksA* and *gluQ-rs*. Deletion or alteration of the predicted terminator resulted in increased expression of the *lacZ* reporter compared with cells containing the wild type terminator sequence. Analysis of the phenotype of a *gluQ-rs* mutant suggested that it may play a role in some stress responses, since growth of the mutant was impaired in the presence of osmolytes.

Conclusions: The results presented here, show that the expression of *gluQ-rs* depends on the *dksA* promoter, and strongly suggest the presence and the functionality of a transcriptional terminator regulating its expression. Also, the results indicate a link between glutamyl-queuosine synthesis and stress response in *Shigella flexneri*.

Keywords: tRNA modification, Gene expression, Stringent response, Osmotic stress

Background

The fidelity of the translation process depends on the aminoacyl-tRNA synthetase enzymes (aaRS). These essential enzymes are responsible for the correct attachment of the corresponding amino acid onto the cognate tRNA, therefore organisms have at least 20 synthetases [1]. The enzymes are divided in two classes, each class having a conserved structure. The genes encoding the aaRS are easily detected within sequenced genomes [2,3], and some species contain synthetase gene

duplications, such as the glutamyl-tRNA synthetases (GluRS) in *Acidithiobacillus ferrooxidans* and *Helicobacter pylori* (genes *gltX1* and *gltX2*) [4,5]. aaRS paralogs, predicted sequences with homology to fragments of synthetases, have also been identified, which is not surprising given the modular nature of the aaRS [6]. Some of the paralogs may be pseudogenes while others have known functions. For instance HisZ from *Lactococcus lactis*, which has similarity with the catalytic domain of histidyl-tRNA synthetase, is involved in histidine biosynthesis [7]. A recent study in *Salmonella enterica* has shown that PoxA, encoded by *poxA/genX*, has similarity to the carboxy-terminal catalytic domain of lysine-tRNA synthetase and is required for posttranslational

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aminoacylation of bacterial elongation factor P. A *poxA* mutant has reduced colonization and virulence, possibly due to misregulated expression of proteins encoded by the SPI-1 pathogenicity island [8,9].

An *Escherichia coli* glutamyl-tRNA synthetase paralog, glutamyl queuosine-tRNA^{Asp} synthetase (GluQ-RS) has approximately 35% amino acid similarity with the catalytic domain of GluRS. This includes the amino acids involved in recognition and activation of glutamate. Although GluQ-RS is missing the carboxyl-terminus domain responsible for the tRNA recognition, in *E. coli* this enzyme is able to activate the amino acid in the absence of the tRNA. Further, once the aminoacyl-adenylate has been formed, the enzyme attaches the glutamate to the nucleoside queuosine present onto the tRNA^{Asp}. Therefore, this enzyme is involved in the synthesis of a new modified nucleoside glutamyl-queuosine (GluQ) present in tRNA^{Asp} [10,11]. This modification is present in tRNA isolated under acidic conditions from bacterial cells grown in rich media. However, the enzyme is not essential for growth of *E. coli* in rich or minimal media [10]. Queuosine is widely distributed in bacteria, and it is present in the first base of the anticodon of tRNA^{Asp}, tRNA^{Asn}, tRNA^{His} and tRNA^{Tyr} [12]; however in *E. coli* only tRNA^{Asp} is a substrate for the GluQ-RS enzyme.

The presence of modifications within the anticodon loop of the tRNA, could enhance the accuracy of the codon binding [13]. Then the tRNA^{AspQ³⁴ might improve recognition of both GAC and GAU codons [14] and stimulate the binding of the GAU codon to the ribosome [15]. In *Shigella flexneri* it has been shown that mutations in genes required for tRNA modifications, *miaA* and *tgt* decreased virulence. *miaA* is required for 2-methylthio-N⁶-isopentenyladenosine modification at position 37 of the anticodon loop and *tgt* is involved in queuosine modification at position 34 within the anticodon loop [16-18]. In this study, we determined the role of the genome organization and its effect on the expression of the *gluQ-rs* gene in the major human pathogen, *S. flexneri*.}

Results

Genomic organization of the *S. flexneri* *gluQ-rs* gene

GluQ-RS is required for the synthesis of the modified nucleoside, GluQ, present on tRNA^{Asp} [10,11]. By searching the bacterial protein database Uniprot (<http://www.uniprot.org/>), we were able to identify GluQ-RS in more than a hundred bacterial species, primarily proteobacteria (Figure 1, filled symbols). From the phylogenetic analysis we can distinguish the three subgroups of enzymes described by Dubois et al., 2004 [11], which are characterized by the presence of the signature HXGS, HXGN or HXGH in the adenylate binding site. A similar tree was obtained using the Neighbor joining method.

Phylogenetic analysis within the subgroup of enzymes with the HXGN motif, included representatives from the Firmicutes bacterial group (Figure 1, open square) together with *Desulfovibrio vulgaris* and *Truepera radiovictrix* enzymes. From the alignment, these members have 8 characteristic amino acids, G₇₀PDXGGXX, that do not align with the other GluQ-RS (Figure 1, numbering is derived from *D. vulgaris* enzyme). Further genomic analysis indicated that the *gluQ-rs* gene is found primarily in two genomic arrangements, either alone or located immediately downstream of *dksA*. Searching within the String database [19] and GenomeNet [20], we found that the *dksA-gluQ-rs* gene organization was conserved in more than 40 different species, all of which were within the gammaproteobacteria group. These included species of *Aeromonadales*, *Alteromonadales*, *Enterobacteriaceae*, including *E. coli* and *S. flexneri*, *Pseudomonadales*, and *Vibrionaceae* (Figure 1).

A bioinformatics analysis of the intergenic region between *dksA* and *gluQ-rs* showed great variation in the distance between the two genes among these bacterial species. In *S. flexneri* the intergenic region between the stop codon of *dksA* and the first codon of *gluQ-rs* is only 39 base pairs. Therefore, we suspected that the transcription of *gluQ-rs* was regulated by the previously characterized *dksA* promoter [22]. To test this hypothesis, we isolated total mRNA and performed RT-PCR to identify an mRNA that included both genes (Figure 2A). The observation that there is an mRNA species containing both genes (Figure 2A, lane 1) indicates that they are co-transcribed and that the expression of *gluQ-rs* may be regulated by the *dksA* promoter.

S. flexneri *gluQ-rs* gene is co-transcribed with *dksA* gene

Although *S. flexneri* *gluQ-rs* can be transcribed from the *dksA* promoter, this did not rule out the presence of an additional, independent promoter. Therefore, the expression of each gene was measured by RT-PCR during different stages of *S. flexneri* growth in Luria Bertani (LB) at pH 7.4. The analysis of the *dksA* and *gluQ-rs* transcripts shows that for both mRNAs, the level is stable during the growth curve, with an increase of 1.3-fold at stationary phase compared to the early mid log phase (Figure 2B, compare lanes 2 and 4). However, the mRNA that includes the intergenic region showed variation depending on the stage of growth, increasing 20-fold at stationary phase compared with its expression at early mid log phase (Figure 2B *dksA/gluQ-rs*, compare lanes 2 and 4). In order to confirm those results, a transcriptional fusion strategy was used. Different segments of the operon were cloned and fused to the *lacZ* reporter gene in pQF50, and promoter activity was assayed by β-galactosidase activity [23]. Kang and Craig, 1990 [22] identified three promoters for *dksA*. By mean of

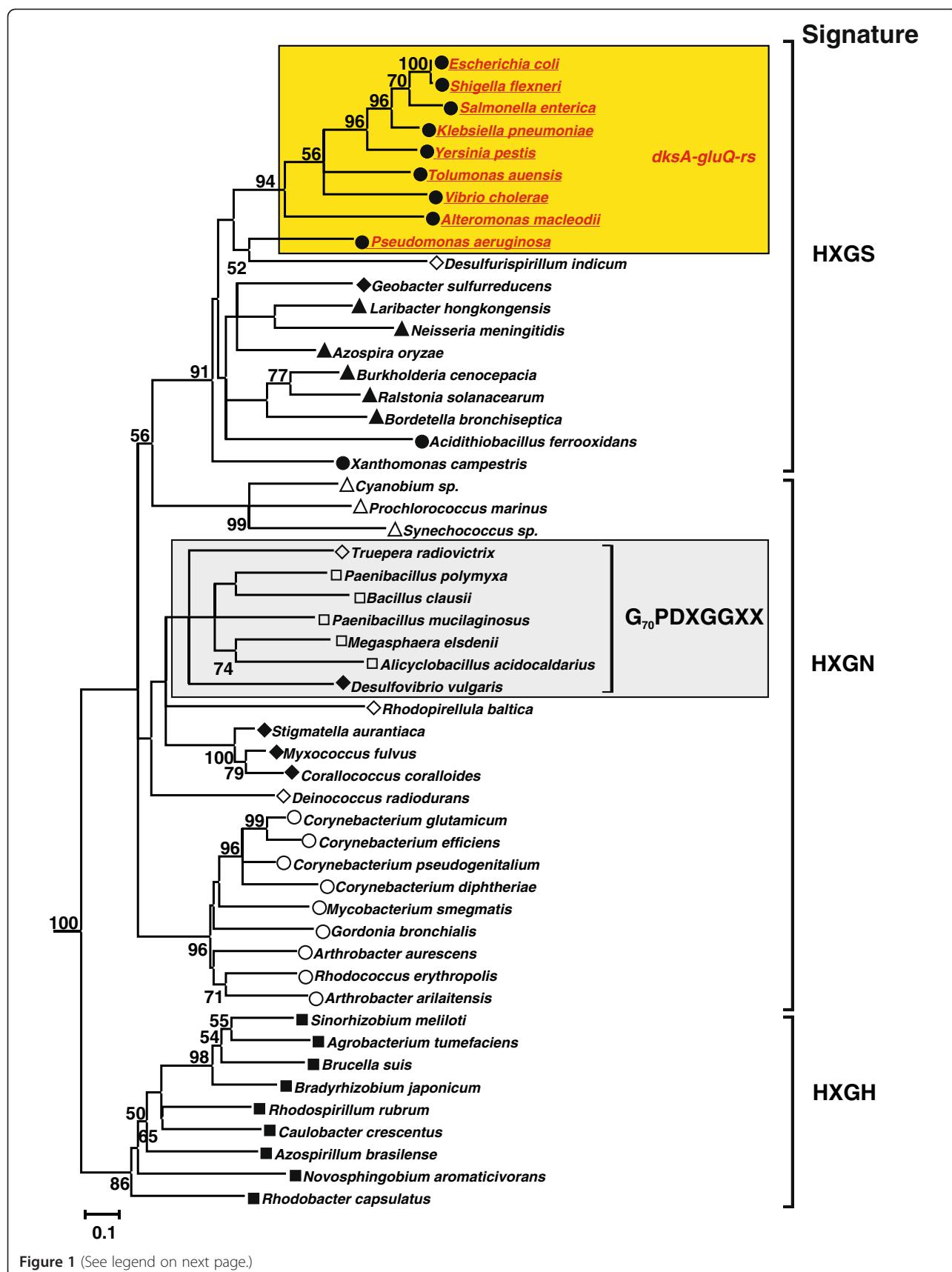


Figure 1 (See legend on next page.)

(See figure on previous page.)

Figure 1 GluQ-RS is distributed within the bacterial domain. Rooted Phylogenetic analysis of selected sequences of GluQ-RS, showing the presence of this enzyme in the bacterial domain. Searching within the Uniprot database (<http://www.uniprot.org/>) of homologues protein to GluQ-RS were identified and then were searched in the GenomeNet (<http://www.genome.jp/>) database for confirmation and analysis of the genomic organization. Bootstrap values (>50%) were calculated by 400 replicates using the maximum-likelihood methods in the MEGA5 software [21] and rooted with archaeal GluRS from *Methanocaldococcus jannaschii* and *Archaeoglobus fulgidus* (not shown). In yellow background are shown bacterial species (in red and underlined) that are representatives having the genomic organization of *dksA-gluQ-rs* genes. The signature of each subgroup identified previously [11] is indicated. Filled symbols representing proteobacteria groups, open symbols represent other bacterial groups. ■: alphaproteobacteria, ▲: betaproteobacteria, ●: gammaproteobacteria, ♦: deltaproteobacteria, O: actinobacteria, △: cyanobacteria, □: firmicutes, ♢: others.

bioinformatics tools, including BPROM from the Softberry software package (<http://linux1.softberry.com/berry.phtml>), we identified those promoters in *S. flexneri* and included all three promoters in the constructs indicated in Figure 3A. The plasmid containing a fragment from the *dksA* promoters to the beginning of the *gluQ-rs* gene, with the first five amino acids of GluQ-RS, named pVCPDT, represents the full length *dksA* gene with its native promoters (Table 1 and Figure 3A). A second fusion construct, pVCDT, contains sequence from the beginning of the coding region of *dksA* through the beginning of *gluQ-rs* and also included the first five amino acids of GluQ-RS. Because pVCDT does not have the *dksA* promoter region, it served as the reporter for transcription from an independent *gluQ-rs* promoter. A third construct, pVCPD, contained the segment from the *dksA* promoter to the end of the *dksA* gene, hence this plasmid does not have the intergenic region, nor the first amino acids of GluQ-RS (Table 1). Each of the recombinant plasmids was transformed into *S. flexneri* and the β-galactosidase activity was measured when the bacterial cells reached mid-log phase. Analysis of the enzymatic activity of these reporter fusions showed that the strain carrying pVCDT had baseline levels of the enzyme (Figure 3B), indicating that there is not an independent promoter for *gluQ-rs*. Thus, the promoter upstream of *dksA* is responsible for the expression of both genes, at least under the conditions assayed (see lane pVCPDT Figure 3B). Therefore, these two results (Figure 2 and Figure 3B) indicate that *dksA* and *gluQ-rs* form an operon, and *gluQ-rs* lacks an additional, separate promoter. A surprising observation was obtained with the clone containing pVCPD, which showed a ten-fold increase in enzymatic activity compared to pVCPDT (Figure 3B). This suggested the presence of a terminator or other regulatory sequence in the intergenic region that modulated the expression of *gluQ-rs*.

The *S. flexneri* *gluQ-rs* gene has an upstream transcription terminator

In order to explain the difference observed in expression of *lacZ* from the recombinant plasmids pVCPDT and

pVCPD a bioinformatic analysis using mFold [26] was performed to search for possible secondary structures in the mRNA. A potential transcriptional terminator was found at the beginning of the *gluQ-rs* gene, leaving the first predicted AUG codon located on the bulge of this terminator (Figure 4A). In order to determine the functionality of this terminator, we performed site directed mutagenesis to disrupt the structure in the predicted stem (Figure 4A). As shown in Figure 4B, the plasmid containing the mutations, pVCPDTMut had >2-fold higher enzymatic activity ($p < 0.05$) than the plasmid containing the wild type sequence. This result suggested that the intergenic region upstream of *gluQ-rs* contains a transcriptional terminator.

Identification of the first methionine

The first methionine in the predicted GluQ-RS protein corresponds to the one located on the bulge of the terminator structure (Figure 4A), which also contains a possible Shine-Dalgarno sequence. However, in related species like *Escherichia fergusonii* that also have the terminator structure, a methionine is not present at that location. In the *S. flexneri* sequence, there is another AUG codon in the same reading frame 27 nucleotides downstream from the one in the terminator. In order to determine which methionine is the start site for translation of the *S. flexneri* GluQ-RS, we constructed a vector that included the intergenic region from the stop codon of the *dksA* gene to the end of *gluQ-rs* cloned into the expression vector pET15c. This allowed expression of C-terminal His-tagged GluQ-RS under T7 promoter control. The protein was partially purified by affinity chromatography as described elsewhere [10], and the sequence of the amino terminus of the GluQ-RS protein was determined to be NH₂-T-D-T-Q-Y-I-G-R-F-A-P, which corresponds to the amino acid sequence after the second methionine. Therefore, the initiator methionine is not the one indicated in the database, and the protein is 298 amino acids. Surprisingly, there is no obvious Shine-Dalgarno sequence adjacent to the initiator methionine we identified (Figure 5).

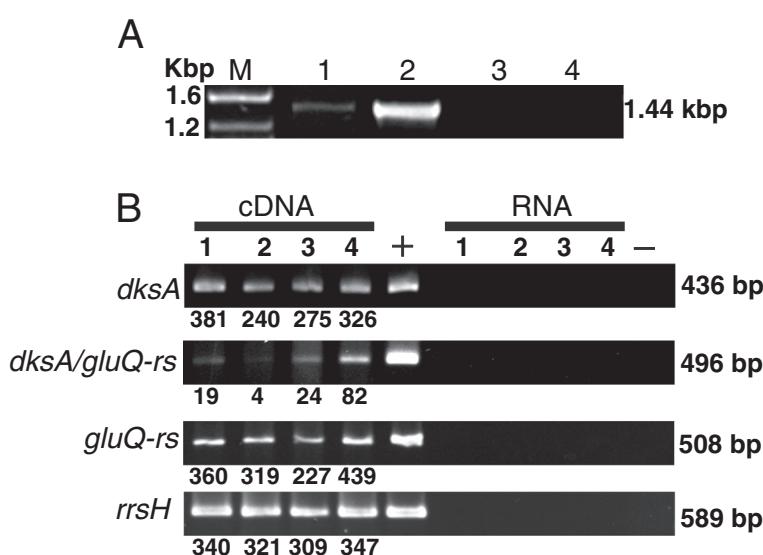


Figure 2 *gluQ-rs* is co-transcribed with *dksA* in *S. flexneri* 2457T. **A)** Agarose gel showing the amplified product of the full-length operon extending from the *dksA* gene through the end of *gluQ-rs* (1.44 kbp). Total RNA isolated during mid log phase growth of *S. flexneri* was subjected to reverse transcriptase and PCR (RT-PCR) using primers opeF/opeR (Table 2). **M:** molecular marker, sizes are indicated. **Lane 1:** RNA treated with reverse transcriptase, **Lane 2:** genomic DNA isolated from *S. flexneri* 2457T, **Lane 3:** RNA without reverse transcriptase treatment, **Lane 4:** negative control of PCR reaction without DNA. **B)** Electrophoretic analysis of each amplified gene fragment, *dksA* (*dksAF/dksAR*; 436 bp), *gluQ-rs* (*gQRSF/gQRSR*; 508 bp), the intergenic region *dksA/gluQ-rs* (*interF/interR*; 496 bp) and the ribosomal RNA 16S (*rrsHF/rrsHR*, 589 bp). Total RNA isolated during different phases of the growth curve of *S. flexneri* 2457T was subjected to RT-PCR to detect the corresponding fragment. **Lane 1:** lag phase, **Lane 2:** early mid log phase, **Lane 3:** mid log phase, **Lane 4:** stationary phase. +: corresponds to amplification using genomic DNA. **RNA:** Isolated RNA without reverse transcriptase treatment. -: negative control PCR reaction without DNA. Each band was estimated using Image J software (V1.46) and its amount was estimated using the fragment corresponding to 500 bp of the DNA ladder.

Phenotype of the *S. flexneri* *gluQ-rs* mutant

To determine the role of GluQ-RS in *S. flexneri* growth and virulence, a deletion mutant of the *gluQ-rs* gene was constructed in *S. flexneri* 2457T. The mutant was compared to the wild type by Biolog phenotype MicroArrays (Biolog, Inc., Almeda, CA). The major difference observed for the mutant was impaired metabolism when grown under osmotic stress conditions (Figure 6). The mutant had a longer lag and reduced growth compared to the wild type in the presence of increasing concentrations of potassium chloride, sodium sulfate, sodium formate, sodium benzoate, sodium nitrate and sodium nitrite. The phenotype was complemented with the *gluQ-rs* gene cloned into an expression vector. No differences were observed in the growth or metabolism of these strains when they were incubated in presence of 1% sodium chloride, which was similar to LB (Figure 6 and data not shown).

Because expression of *dksA* is required for *S. flexneri* virulence [27], and growth of *Shigella* in the intracellular environment may induce a stress response, we also measured invasion and plaque formation by the *gluQ-rs* mutant. However, no significant differences were noted (data not shown), suggesting that GluQ-RS is not essential for invasion or intracellular growth of *S. flexneri*.

Discussion

Conserved *dksA-gluQ-rs* genomic organization in gammaproteobacteria

GluQ-RS, a paralog of GluRS synthetase, is involved in the formation of GluQ, the nucleoside located at the wobble position of tRNA^{Asp} in bacteria. The protein is present in Firmicutes, Actinobacteria, Cyanobacteria, Alphaproteobacteria, Betaproteobacteria, Gammaproteobacteria and Deltaproteobacteria (Figure 1). From the phylogenetic analysis we distinguished the three subgroups described previously [11] based on the HIGH motif that is present in the class I aminoacyl-tRNA synthetases [2]. As was described previously [11], all GluQ-RS enzymes are characterized by the replacement of a threonine in GluRS enzymes, which is involved in the recognition of the amino acid and the terminal adenosine of the tRNA^{Glu} (Thr₁₃₃ of *Methanocaldococcus jannaschii* GluRS enzyme) by isoleucine, leucine or valine at that position (Ile₄₇ of *S. flexneri* GluQ-RS). This substitution is also conserved in all enzymes analyzed here, including those from the Firmicutes group.

The *gluQ-rs* gene is widely distributed in the bacterial domain; however, its genome organization is variable. We observed that only in members of the gammaproteobacteria, namely *Aeromonadales*, *Alteromonadales*, *Pseudomonadaceae*,

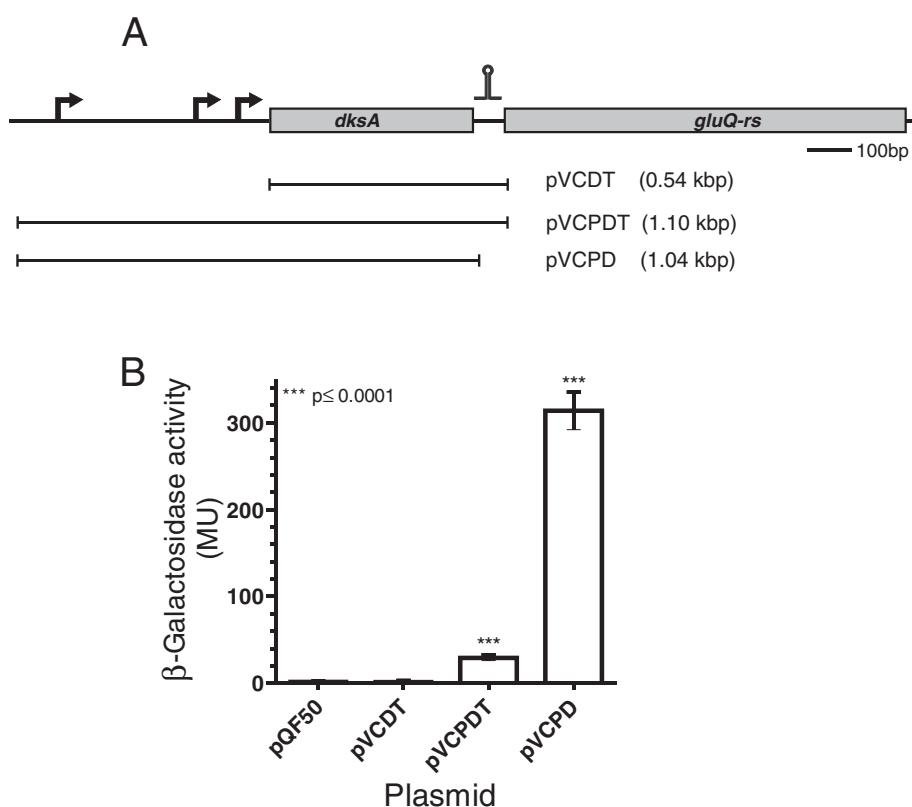


Figure 3 The transcription of the *gluQ-rs* gene is controlled by a termination stem loop. **A**) Schematic representation of the operon, the arrows indicate the position of each promoter identified by our bioinformatics analysis and experimentally determined by Kang and Craig, 1990 [22]. The putative ρ -independent terminator is represented by the stem loop symbol upstream of *gluQ-rs* gene. The horizontal bar represents the DNA region amplified and cloned into pQF50 (Table 1). The recombinant plasmids are described in Table 1. **pVCDT** does not have the *dksA* promoter but has the terminator. **pVCPT** has the promoter region of *dksA* and the terminator upstream of *gluQ-rs*; therefore, it represents the genomic organization of the operon. **pVCPD** also has the promoter of *dksA* but lacks the terminator region. The size of each fragment is indicated. **B**) β -galactosidase activity of each protein extract obtained from the corresponding clone. The data represent the average of three experiments in triplicates and the Student *t* test was used to compare the means between each clone with the empty vector. *** *p* values <0.05 were considered statistically significant.

Enterobacteriaceae and *Vibrionaceae*, the *gluQ-rs* gene is located immediately downstream of the *dksA* gene (Figure 1). A more detailed analysis shows that even within this genomic organization there are differences. In some species of *Pseudomonadaceae*, such as *P. aeruginosa*, *P. entomophila*, and *P. fluorescens*, we observed the same genomic structure as in *E. coli* or *S. flexneri*, with a distinctive terminator between the genes. In contrast, while the *dksA* gene is also upstream of *gluQ-rs* in some *P. syringae*, there are insertions of an encoded transposase or more than a 400 base pairs separating both genes without a detectable terminator. However, using bioinformatics tools we detected a possible promoter within this region in *P. syringae* (data not shown), indicating that the expression of the *gluQ-rs* gene may be under control of its own promoter, a question that remains to be addressed.

Stringent response and tRNA modification

Our bioinformatic analysis shows the presence of a transcriptional terminator and lack of a *gluQ-rs*-specific

promoter. This is consistent with our results, in which we did not detect any activity from promoters other than those upstream of the *dksA* gene (Figure 3). This unusual arrangement suggests that *gluQ-rs* expression is dependent on *dksA*-regulated conditions. Because DksA is a key member of the stringent response in bacteria and regulates a number of processes in the cell, including its own expression [25,28], the data suggest that there is coordinate regulation of tRNA modification and other DksA targets. Although we could not detect any promoter activity specific for *gluQ-rs* in the growth conditions tested (i.e. altering the pH, presence of glutamate), we cannot discount the possibility that the gene is specifically regulated under some other conditions. The regulon database (<http://regulondb.ccg.unam.mx/index.jsp>) indicates that the *E. coli* *gluQ-rs* gene has a recognition site for the σ^{24} subunit of RNA polymerase. From our analysis, this sequence is identical to *S. flexneri*, but there is no experimental evidence of this recognition. Interestingly, when the *gluQ-rs* gene was

Table 1 Bacterial strains and plasmids used in this work

Bacterial strains or plasmid	Characteristics	Source or reference
<i>Shigella flexneri</i>		
<i>S. flexneri</i> 2457T	Wild type strain	Laboratory stock
<i>S. flexneri</i> 2457T Δ gluQ-rs::kan	Deletion mutant of gluQ-rs gene	This work
<i>Escherichia coli</i>		
<i>E. coli</i> W3110 Δ gluQ-rs::kan	Deletion mutant of gluQ-rs gene	[10]
DH5a	F^+ $\phi 80lacZ\Delta M15$ $\Delta(lacZYA-argF)$ $U169$ $recA1$ $endA1$ $hsdR17$ (rK , mK +) $phoA$ $supE44$ λ - $thi-1$ $gyrA$ $relA1$	[24]
BL21(DE3)	F^+ $ompT$ gal dcm lon $hsdS_B(r_B\ m_B)$ λ (DE3) [$lacI$ $lacUV5-T7$ gene 1 $ind1$ $sam7$ $nin5$])	Invitrogen
Plasmids		
pTZ57R/T	<i>bla</i> , pMB1 ori, <i>lacZ</i> peptide, f1 phage ori	Fermentas®
pQF50	<i>bla</i> , pMB1 ori, <i>lacZ</i> gene without promoter	[23]
pET15c	Empty vector, a modified version of pET15b	This work
pVCDT	<i>S. flexneri</i> fragment from nucleotide +58 ^a of <i>dksA</i> gene to beginning of <i>gluQ-rs</i> gene (+590) cloned into pQF50. Pair of primers used were PgluQF/PdksARCT.	This work
pVCPDT	<i>S. flexneri</i> fragment from nucleotide -506 of <i>dksA</i> gene to beginning of <i>gluQ-rs</i> gene (+590) cloned into pQF50. Pair of primers used were PdksAF/PdksARCT.	This work
pVCPDMut	<i>S. flexneri</i> fragment from nucleotide -506 of <i>dksA</i> gene to beginning of <i>gluQ-rs</i> gene (+590) cloned into pQF50, with the terminator mutated by the nucleotides indicated in Figure 4a.	This work
pVCPD	<i>S. flexneri</i> fragment from nucleotide -506 of <i>dksA</i> gene to nucleotide +527 (end of <i>dksA</i> gene) cloned into pQF50. Pair of primers used were PdksAF/PdksARST.	This work
pATGGQRS	<i>S. flexneri</i> gene from nucleotide +509 (stop codon of <i>dksA</i>) to nucleotide +1469 (last codon of <i>gluQ-rs</i> without stop codon). Pair of primers used were ATGGQRSF/ATGGQRSR.	This work

^aFragments cloned are numbered based on the transcription start of *dksA* identified by [25].

deleted in *S. flexneri*, the mutant showed impaired growth in the presence of osmolytes (Figure 6). A recent publication demonstrated that σ^{24} and σ^S proteins from *Salmonella enterica* serovar Typhi are important for the expression of several genes induced by osmotic stress in this bacterium [29]. Moreover, the expression of the gene encoding σ^{24} in *E. coli* is regulated by the stringent response [30]. The possible role of σ^{24} on the expression of *gluQ-rs* under osmotic stress might be interesting to study.

GluQ-RS is an enzyme responsible for the formation of the GluQ tRNA modification, and two independent groups [10,11] have shown that this enzyme required a high concentration of glutamate to be activated and transferred to the queuosine base present on the tRNA^{ASP}. Interestingly, one of the first events to occur when bacteria are subject to high osmolyte stress is an increase in glutamate levels within the cytoplasm [31]. Our observation indicates an important role of the tRNA modification for the growth of *S. flexneri* in the presence of osmolytes (Figure 6). Other tRNA modifications might play a similar role in this stress condition. In *E.*

coli, inactivation of the *yfiC* gene, responsible for the modification at the adenosine 37 present on the tRNA^{Val}, leads to a high sensitivity to osmotic stress [32].

Transcription of *gluQ-rs* is regulated by a terminator

The results obtained in the present work show the presence of a terminator and suggested the functionality of this structure (Figure 3 and Figure 4). To our knowledge, there are few examples of bacterial genes that have similar structures. There is a terminator structure upstream of the DNA primase gene, *dnaG*, which also has an unusual Shine Dalgarno sequence [33]. Another example is the *recX* gene in *E. coli*, where readthrough accounts for approximately 10% of its transcription [34].

The two characteristics of *gluQ-rs* described in this work, co-transcription with the upstream gene and the presence of a terminator immediately upstream, allow us to propose that both the transcription and translation process could be regulated in the *gluQ-rs* gene. It has been described, that the presence of terminators upstream of the coding region might be part of a regulatory system such as a riboswitch [35]. Riboswitches for

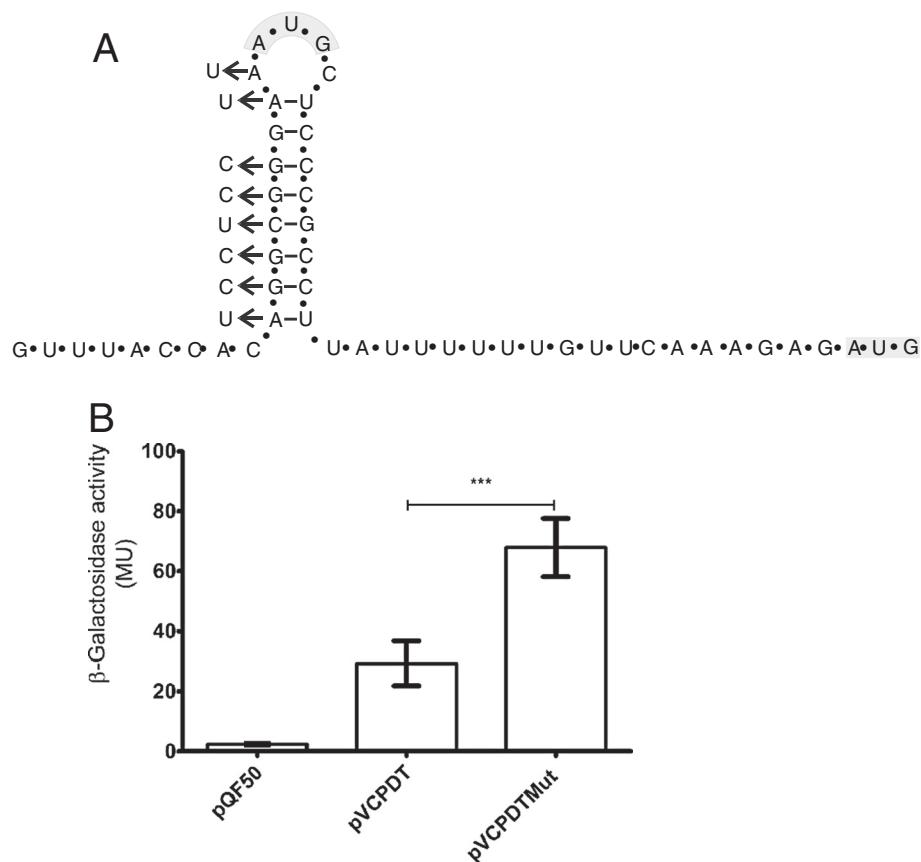


Figure 4 Functionality of the transcriptional terminator upstream of *gluQ-rs*. **A)** Schematic representation of the terminator with $\Delta G = -14.7$ Kcal/mol identified using Mfold software [26]. Bases shaded in grey indicate the two possible AUG start codons, one located in the bulge of the terminator structure and the other located 27 nucleotides downstream. The arrows indicate the site directed mutagenesis location, with the corresponding nucleotide changes designed to disrupt the predicted structure. **B)** β -galactosidase activity of protein extracts obtained from the corresponding clones. The plasmid **pVCPDTMut** has a similar construction as **pVCPDT** but contains the mutated terminator indicated above. The data represent the average of three experiments, each done in triplicate, and the Student *t* test was used to compare means between the pVCPDT and pVCPDTMut clones. *** *p* values <0.05 were considered statistically significant.

genes involved in queuosine formation have been described, in which the precursor preQ1 is the ligand of the mRNA structure [36]. Using the riboswitch server (ribosw.mbc.nctu.edu.tw [37]), we did not identify any potential riboswitch (data not shown). However we cannot discount that the terminator described here might be part of a regulatory circuit similar to a riboswitch, or that an unidentified protein might bind the terminator structure.

GluQ modification and codon bias

tRNA modifications present at the anticodon loop might be important for the accuracy of codon reading during the translation processes [13]. Morris et al., 1999 [14] proposed, based on molecular modeling, that the tRNA^{AspQ34} might improve recognition of both GAC and GAU codons, consequently the interaction of the codon GAU with the anticodon of tRNA^{AspG34} could be less efficient. In fact, in *S. flexneri* there are a few genes

such as *sitA*, *virF* and *proX* (an inducible gene under osmotic stress) that have a bias toward those codons that favor the modified tRNA. Thus, while there is no obvious loss of plaque formation in the *gluQ-rs* mutant (data not shown), the absence of GluQ-RS may influence the expression of proteins such as Sita that are required for fitness of *Shigella* in the host [38].

Conclusions

In this work we have shown that the expression of *gluQ-rs*, a gene codifying an enzyme involved in the formation of GluQ present on the tRNA^{Asp}, is under the control of the *dksA* promoter. Also, we show the presence of a functional terminator that controls the expression of *gluQ-rs*. Finally, we present data that suggest that the presence of modification of the tRNA^{Asp} is important for survival of the human pathogen *Shigella flexneri* under osmotic stress conditions.

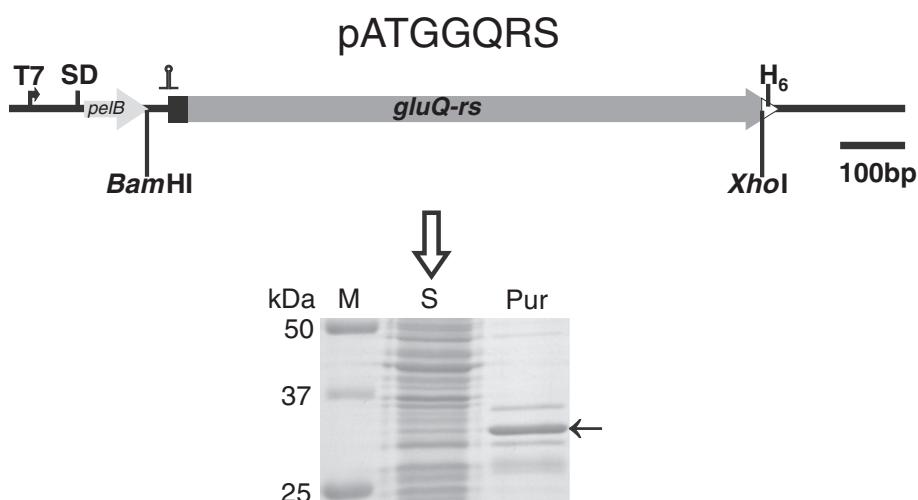


Figure 5 Determination of the first methionine of GluQ-RS. The cloning strategy utilized is shown at the top. A fragment from the stop codon of *dksA* to the end of *gluQ-rs* gene was amplified from *S. flexneri* genomic DNA with the primers ATGGQRSF/ATGGQSR and cloned into a pET15c vector using the restriction sites *Bam*HI and *Xba*II. Therefore this clone represents the operon, but *dksA* was replaced with the plasmid encoded fragment *pelB*. The transcription of this plasmid, named pATGGQRS, is controlled by the T7 promoter and translation is controlled by the Shine-Dalgarno (SD) sequence of *pelB*, both contained within the plasmid. The GluQ-RS protein synthesized has a histidine tag (*H₆*) at the C-terminus, facilitating protein purification. The putative p-independent terminator is represented by the stem loop symbol upstream of *gluQ-rs* gene, and the black box in the gene indicates the two possible peptides, depending of which methionine is utilized. Bottom: The SDS -polacrylamide gel electrophoresis showing the supernatant extract (S) and the partially purified protein (Pur) produced by cells carrying the recombinant plasmid. The predominant band indicated by the arrow was excised and subjected to amino terminal sequencing, yielding the following amino acid sequence: T-D-T-T-Q-Y-I-G-R-F-A-P. This corresponds to the sequence following the second methionine. The size of the molecular markers (M) are given in kDa.

Methods

Bacterial growth conditions

The bacterial strains and plasmids used in this study are described in Table 1. *E. coli* strains were maintained on LB-agar (10 g of tryptone per liter, 5 g of yeast extract per liter, 10 g of NaCl per liter and 15 g of agar per liter), *Shigella* strains were maintained on Trypticase Soy Agar plus 0.01% congo red. All strains were stored at -80°C in LB broth plus 20% glycerol. The bacteria were grown in LB broth adjusted

to pH 7.4 with 40 mM MOPS (Merck) or M9 minimal media [24]. When necessary, ampicillin was added to a final concentration of 100 µg/ml. Bacterial growth was monitored by optical density at 600 nm (OD₆₀₀).

Bioinformatics tools to construct the phylogenetic tree

The protein sequences were obtained from the Uniprot database (<http://www.uniprot.org/>) and then were searched in the GenomeNet (<http://www.genome.jp/>) to confirm the

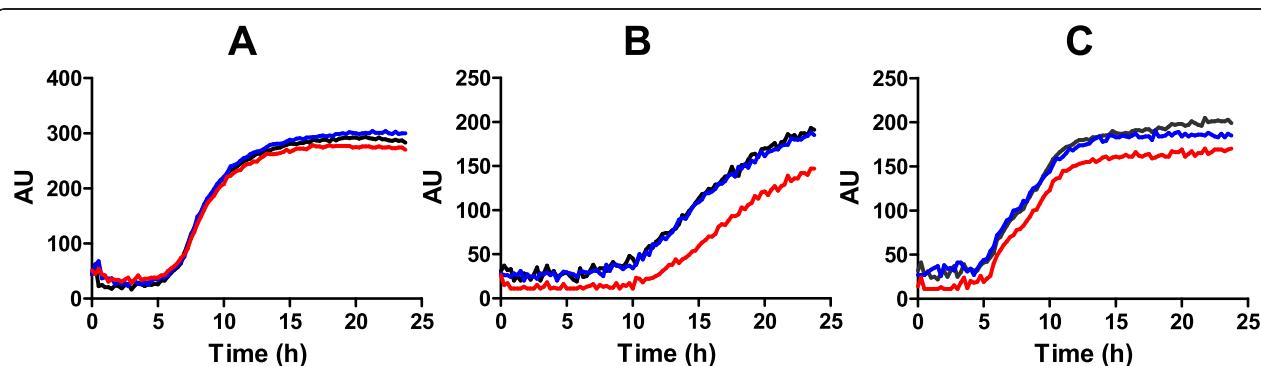


Figure 6 The *gluQ-rs* mutant is sensitive to growth in high osmolarity. Biolog phenotypic MicroArrays were used to characterize the growth and metabolism of the *gluQ-rs* mutant and its wild type parent, 2457T. Wild type (black line) transformed with the empty plasmid, 2457T Δ*gluQ-rs*:kan (red line) transformed with the empty plasmid pCM and *S. flexneri* Δ*gluQ-rs*:kan (blue line) transformed with the complementary plasmid pTRCGQ were incubated for 24 h with medium in the corresponding plate for osmolytes (PM9). **A)** sodium chloride 1% **B)** sodium benzoate 20 mM pH 5.2. **C)** sodium nitrate 100 mM. Metabolism was monitored by measuring reduction of the tetrazolium dye in the medium at 15 min intervals and is shown as units.

genomic organization. A selected number of GluQ-RS enzymes were aligned using the MUSCLE algorithm [39] and analyzed using the maximum-likelihood method based on the JTT matrix-based model. The percentage of trees in which the associated proteins clustered together is shown next to the branches. The analysis involved 54 amino acid sequences, including the GluRS proteins from *Methanocaldococcus jannaschii* and *Archaeoglobus fulgidus* as an outgroup. All positions containing gaps and missing data were eliminated. There were a total of 199 positions in the final dataset. Evolutionary analyses were conducted in MEGA5 [21].

RNA isolation and synthesis of cDNA

Total mRNA was obtained during the growth of *S. flexneri* 2457T using the RNeasy mini kit following the supplier instructions (Qiagen). The purified nucleic acid was treated with RNase- free DNase (Fermentas) and its concentration was estimated by measuring the optical density at 260 nm (OD₂₆₀). Approximately 1 μg of total RNA was subjected to reverse transcription using M-MuLV polymerase (Fermentas) and random primers following the

provider's protocol. The cDNA was amplified using specific PCR primers for each gene of interest (Table 2).

Construction of transcriptional fusions

Transcriptional fusions were constructed to study the expression control of *gluQ-rs*. Fragments of the *dksA-gluQ-rs* region were fused to *lacZ* in the vector pQF50 by using the *Bam*HI and *Hind*III restriction sites [23]. Each fragment was amplified from *S. flexneri* genomic DNA using the indicated primers (Tables 1 and 2) with the High Fidelity PCR Enzyme Mix polymerase (Fermentas) and cloned into pQF50 (Table 1). Once the sequence of each clone was confirmed, the recombinant plasmid was introduced into *S. flexneri* 2457T by electroporation. The nomenclature of the recombinants plasmids is: P for promoter of the *dksA* gene, D for the *dksA* gene and T for a terminator structure.

β-galactosidase activity

S. flexneri transformed with the corresponding constructs were cultured overnight in LB, a 1:50 dilution was inoculated into 10 ml culture of LB pH 7.4 and grown to an OD₆₀₀ of 0.5. Aliquots of 0.5 ml of each strain containing the clone or the empty vector were

Table 2 Primer sequences

Name	Sequence 5'- 3' ^a	Reference and characteristics
opeF	TAAGGAGAACATGCAAGA	This work. RT-PCR of <i>dksA</i> operon from nucleotide +40 to +147 ^b
opeR	ATAGCTCAGCATGACGCATT	
dksAF	ATGCAAGAAGGGCAAAACCG	This work. RT-PCR of <i>dksA</i> gene from nucleotide +54 to +488
dksAR	GCGAATTTCAGCCAGCGTT	
interF	AGTGGAGACGAAGATTTCG	This work, RT-PCR of intergenic region from nucleotide +368 to +863
interR	TCCTTGTTCATGTAACCAGG	
gQRSF	TTCAAAGAGATGACAGACACAG	This work, RT-PCR of <i>gluQ-rs</i> gene from nucleotide +567 to +1074
gQRSR	CACGGCGATGAATGATAAAATC	
rrsHF	CCTACGGGAGGCAGCAG	[40] RT-PCR of ribosomal RNA 16S
rrsHR	CCCCGTCAATTCTTGAGTT	
pcnBR	GATGGAGCCGAAAATGTTGT	Reverse of <i>pcnB</i> gene from nucleotide +1993
PdksAF	GGATCC AAGCGAAGTAAATACGG	<i>Bam</i> HI site, from nucleotide -506
PdksARST	AAGCTT GTGATGGAACGGCTGTAAT	<i>Hind</i> III site, to nucleotide +527
PdksARCT	AAGCTT CTGTGTCTGTCATCTTTG	<i>Hind</i> III site, to nucleotide +590
PgluQF	GGATCC AAGAGGGCAAAACCGTA	<i>Bam</i> HI site, from nucleotide +58
TERGQ2	CCTTATTTTGTCAAAAGAGATGACAGACACACAGA	Recognition from nucleotide +555
TERMGQ3	ATAAGGGGGAGCATA <u>ACGGAGG</u> GTGGTAAAC	Recognition from nucleotide +560, underline sequence are nucleotides changed
M13R	GCGGATAACAATTTCACACAGG	Recognition site in pTZ57R/T
ATGGQRSF	GGATCC <u>GTAATT</u> ACAGCCGTTCCATC	<i>Bam</i> HI site, from nucleotide +507. Underline nucleotides correspond to the stop codon of <i>dksA</i>
ATGGQSR	CTCGAG GCATGACGCATTGAGAATG	<i>Xba</i> I site, to nucleotide +1469
virFF	AGCTCAGGCAATGAAACTTGAC	[41]
virFR	TGGGCTTGTATTCGATAAGTC	

^aNucleotides in bold are indicated restriction site.

^bFragments cloned are indicated based on the transcription start of *dksA* identified by [25].

assayed for β -galactosidase activity according to Miller [42]. The data were analyzed using the software GraphPad Prism V5.01.

Site directed mutagenesis

A possible transcription terminator between *dksA* and *gluQ-rs* was identified using the program Mfold [26]. Site directed mutagenesis by overlap PCR was performed to disrupt the predicted terminator [43]. Using the fragment VCPDT cloned in the vector pTZ57R/T as template, was amplified a 1,072 bp fragment, which include the mutation, using the primers PdksAF and TERMGQ3, while a second fragment of 162 bp overlapping the mutated region, was obtained with primers TERGQ2 and M13R (Table 2). Both fragments (1,072 bp and 162 bp) were digested with *Dpn*I, purified and mixed at equimolar quantities to carry out a PCR reaction using the 5' and 3' ends primers (PdksAF and PdksARCT). The 1,110 bp amplified fragment was cloned in the vector pTZ57R/T and sequenced to verify the mutation. This plasmid was digested with *Bam*HI and *Hind*III and the fragment subcloned in to the vector pQF50.

Determination of first methionine of GluQ-RS

In order to establish which is the first AUG codon of *gluQ-rs*, the recombinant plasmid pATGGQRS was constructed. A PCR reaction was performed using the primers ATGGQRSF and ATGGQRSR (Table 2) and genomic DNA from *S. flexneri*. The amplified fragment, containing the *Bam*HI site, stop codon of *dksA*, the intergenic region with the terminator, the *gluQ-rs* reading frame without its stop codon and the *Xba*I site was cloned into pET15c, a modified version of pET15b, which was constructed by inserting the 290 bp *Xba*I and *Bpu*I fragment of pET20b containing the polylinker into pET15b. This construct allowed the synthesis of a C-terminal histidine tagged protein under the transcription control of the T7 promoter. The construct was transformed in BL21(DE3) strain and the His-tagged protein was partially purified by affinity chromatography as described previously [10]. The eluted protein was transferred to a PVDF membrane and stained with Coomassie blue. The predominant band of the expected size (34.6 kDa) was sequenced at the Protein Core Facility of the Institute for Cellular and Molecular Biology, University of Texas at Austin.

Construction of the plasmid for complementation of the *gluQ-rs* mutation

This plasmid was constructed from the pATGGQRS plasmid in which the T7 promoter was removed by digestion with *Bgl*II and *Nco*I enzymes and replaced by the TRC promoter obtained from pTRC99a plasmid by amplification and digestion with *Bam*HI and *Nco*I to

obtain the pTRCGQ plasmid. The empty plasmid (pCM) was constructed by incorporating the TRC promoter into the pET15c plasmid.

Inactivation of *gluQ-rs* gene in *S. flexneri*

Deletion of *gluQ-rs* was carried out using the λ red recombinase method [44] with the following modifications. *S. flexneri* 2457T carrying pKD46 and prepared as described elsewhere [44] was transformed with a purified PCR fragment amplified from the *E. coli* Δ *gluQ-rs::kan* mutant strain using primers dksAF and pcnBR (Table 2), increasing the homologous DNA region to more than 450 bp at each side. The mutant was isolated following overnight growth at 37°C on LB-agar containing kanamycin (50 μ g/ml). The deletion was confirmed by PCR using the same pair of primers (dksAF-pcnBR) and using each primer together with an internal primer as described previously [44]. The presence of the *S. flexneri* virulence plasmid was also confirmed by PCR amplification of the *virF* gene using primers virFF and virFR (Table 2).

Effect of the absence of *gluQ-rs* gene in *S. flexneri* metabolism

The effect of the deletion of the *gluQ-rs* gene on the metabolism of *S. flexneri* was analyzed by Biolog phenotype MicroArrays following the manufacturer's instructions (Biolog, Inc., Almeda, CA). Strains were grown at 30°C overnight and 5 ml of LB was inoculated with a 1:100 dilution and grown at 37°C to reach an OD_{650nm} of 0.5. The cells were then washed and resuspended to 2.5×10^7 cfu/ml and diluted 200 fold in to a solution of IF-10a medium (Biolog). Each well was inoculated with 1.2×10^4 cfu (0.1 ml per well) into the corresponding plates and incubated for 24 hrs at 37°C. The metabolism was recorded and analyzed by the Omnilog software (V 1.20.02) (Biolog, Inc., Almeda, CA).

Abbreviations

*Gl*Q-RS: Glutamyl queuosine-tRNA^{Asp} synthetase and its codifying gene; tRNA^{Asp}: Transfer ribonucleic acid codifying aspartic acid; *Gl*Q: Glutamyl-queuosine; aaRS: Aminocetyl-tRNA synthetase; *Glu*RS: Glutamyl-tRNA synthetase; Q: Queuosine; SPI-1: Pathogenicity islands of *Salmonella*; *mia*A: tRNA Δ (2)-isopentenylpyrophosphate transferase; *tgt*: tRNA guanine transglycosylase; RT-PCR: Reverse transcription-polymerase chain reaction; LB: Luria Bertani; MOPS: 3-(N-morpholino)propanesulfonic acid; M-MuLV: Moloney Murine Leukemia Virus Reverse Transcriptase; PVDF: Polyvinyl difluoride.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

VCC: performed cloning, the enzymatic assay, data analysis. VPT: performed *ex vivo* assays of the wild type and mutant strains of *Shigella* and the enzymatic activity under different growth conditions. CM: performed the RT-PCR of the mRNA isolated from *Shigella flexneri*. CRF: performed the Biolog assay in collaboration with JCS. JCS: Analysis of the data, experimental design, N terminal determination of GluQ-RS. SMP and JCS: analyzed the

results and wrote the paper. All authors contributed to the editing and approved the final paper.

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